- $\begin{array}{ccc} \hbox{(iii)} & \hbox{Customer} & \hbox{service/Grievance} \\ \hbox{process.} \end{array}$
- (iv) At least 3 years experience in furnishing Part B injectable drugs.
- (v) Financial performance and solvency.
- (vi) Record of integrity and the implementation of internal integrity measures.
 - (vii) Internal financial controls.
- (viii) Acquisition of all CAP drugs directly from the manufacturer or from a distributor that has acquired the products directly from the manufacturer.
- (ix) Maintenance of appropriate licensure to supply CAP drugs in States in which they are supplying CAP drugs.
- (x) Cost-sharing assistance as described in §414.914(g).
- (xi) Other factors as determined by CMS.
- (2) Approved CAP vendors must also meet the contract requirements under §414.914.
- (c) Additional considerations. CMS may refuse to award a contract or terminate an approved CAP vendor contract based upon the following:
- (1) Suspension or revocation by the Federal or State government of the entity's license for distribution of drugs, including controlled substances.
- (2) Exclusion of the entity under section 1128 of the Act from participation in Medicare or other Federal health care programs. These considerations are in addition to CMS' ability to terminate the approved CAP vendor for cause as specified in §414.914(a).
- (3) Past violations or misconduct related to the pricing, marketing, distribution, or handling of drugs provided incident to a physician's service.
- (d) Multiple source drugs. In the case of multiple source drugs, there must be a competition among entities for the acquisition of at least one CAP drug within each billing and payment code within each category for each competitive acquisition area.
- (e) Multiple contracts for a category and area. The number of bidding qualified entities that are awarded a contract for a given category and area may be limited to no fewer than two.
- $[70~{\rm FR}~39094,~{\rm July}~6,~2005,~{\rm as}~{\rm amended}~{\rm at}~70~{\rm FR}~70333,~{\rm Nov}.~21,~2005;~72~{\rm FR}~66402,~{\rm Nov}.~27,~2007;~74~{\rm FR}~62013,~{\rm Nov}.~25,~2009]$

§414.910 Bidding process.

- (a) Entities may bid to furnish CAP drugs in all competitive acquisition areas of the United States, or one or more specific competitive acquisition areas.
- (b) The amount of the bid for any CAP drug for a specific competitive acquisition area must be uniform for all portions of that competitive acquisition area.
- (c) A submitted bid price must include the following:
- (1) All costs related to the delivery of the drug to the participating CAP physician.
- (2) The costs of dispensing (including shipping) of the drug and management fees. The costs related to the administration of the drug or wastage, spillage, or spoilage may not be included.

[70 FR 39095, July 6, 2005]

§414.912 Conflicts of interest

- (a) Approved CAP vendors and applicants that bid to participate in the CAP are subject to the following:
- (1) The conflict of interest standards and requirements of the Federal Acquisition Regulation (FAR) organizational conflict of interest guidance, found under FAR subpart 9.5.
- (2) Those requirements and standards contained in each individual contract awarded to perform functions under section 1847B of the Act.
- (b) Post-award conflicts of interest. Approved CAP vendors must have a code of conduct that establishes policies and procedures for recognizing and resolving conflicts of interest between the approved CAP vendor and any entity, including the Federal Government, with whom it does business. The code of conduct which is submitted as part of the application must—
- (1) State the need for management, employees, contractors, and agents to comply with the approved CAP vendor's code of conduct, and policies and procedures for conflicts of interest; and
- (2) State the approved CAP vendor's expectations for management, employees, contractors, and agents to comply with the approved CAP vendor's code of conduct, and policies and procedures